

```

=>
=> s interferon(a)(alpha or beta) (p) diabetes mellitus
    4210 INTERFERON
    244295 ALPHA
    162655 BETA
    6991 DIABETES
    2514 MELLITUS
    2494 DIABETES MELLITUS
        (DIABETES(W)MELLITUS)
L3      6 INTERFERON(A) (ALPHA OR BETA) (P) DIABETES MELLITUS

=> d 1-6 kwic

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US PAT NO: 5,624,895 :IMAGE AVAILABLE: L3: 1 of 6

SUMMARY:

BSUM(22)

Researchers have disclosed that both gamma interferon and **alpha interferon** expression may be used to induce Type I **diabetes mellitus** in transgenic mice (Cell, 1988, 52, 773 to 782; and Science, 1993, 260, 1942-1946). Transgenic mice which express either of.

SUMMARY:

BSUM(30)

Also, Koivisto et al, Diabetologia, 1984, 27, 193-198, teach that human leukocyte (**alpha interferon**) administration in patients which have been newly diagnosed with Type I **diabetes mellitus**, in conjunction with insulin administration, results in no higher remissions than patients who have been treated by conventional insulin therapy. Furthermore, Fabris et al, Lancet, 1992, 340, 548 recently reported a patient that developed Type I **diabetes mellitus** during leukocyte interferon therapy (for chronic human hepatitis) and hypothesized that this treatment may have enhanced the autoimmune process, although.

DETDESC:

DETD(23)

Other moieties which may be fused to gamma interferon include therapeutic agents which are used for treatment of Type I **diabetes mellitus** e.g., immunosuppressive drugs such as cyclosporin, SK506, azathioprine, CD3 antibodies and steroids. Also, gamma interferon may be fused to immunostimulants, immune modulators, and other cytokines such as alpha or **beta interferon**.

US PAT NO: 5,565,423 :IMAGE AVAILABLE: L3: 2 of 6

SUMMARY:

BSUM(29)

Desmopressin diabetes insipidus

Corticotripin inflammatory 39
(ACTH) diseases

Tetracosactide	inflammatory diseases	24
Alsactide	"	17
Insulin	diabetes mellitus	51
beta-sleep ind.		
Peptide	sleep disturbances	9
Secretin	gastric hemorrhages	27
Cholecystokinin	diseases of the	
	matory disorders	
Atriopeptin III	cardiac and renal	24
	insufficiency	
ANF-(99-126)	"	28
Thymopentin	rheumatoid arthritis	5
Interferon-alpha	colds	125
Thyroliberin	hypophysis diagnostic	3
(TRH)		
Gonadoliberin	cryptorchism, sterility	10
(LHRH)		
Buserelin	prostate cancer,	9.

US PAT NO: 5,534,269 :IMAGE AVAILABLE:

L3: 3 of 6

SUMMARY:

BSUM(136)

Indications . . . C, HBe antigen positive chronic active hepatitis B), cancers (e.g., renal cancer and multiple myeloma) when the water-soluble polypeptide is **interferon alpha**, anemia (e.g., anemia during renal dialysis) when the water-soluble polypeptide is erythropoietin, neutropenia (e.g., during anticancer agent therapy) and infectious. . . is FGF-9, senile dementia and neuropathy when the water-soluble polypeptide is NGF, thrombosis etc. when the water-soluble polypeptide is TPA, **diabetes mellitus** when the water-soluble polypeptide is insulin, and cancers when the water-soluble polypeptide is tumor necrosis factor.

US PAT NO: 5,417,982 :IMAGE AVAILABLE:

L3: 4 of 6

SUMMARY:

BSUM(42)

The . . . to entrap other growth hormones in a polymer matrix, e.g. estrogens, androgens, insulin, IGF, interleukin-I and interleukin-II. Cytokins such as **interferon-beta** and **interferon-gamma**, used in the treatment of diseases such as osteoporosis, **diabetes mellitus**

and multiple sclerosis may also benefit from the present invention.

US PAT NO: 5,165,921 :IMAGE AVAILABLE:

L3: 5 of 6

SUMMARY:

BSUM(7)

In addition, others have treated condyloma acuminata with recombinant human **alpha-interferon** by the subcutaneous and intramuscular injection of interferon. Both recombinant human **alpha-interferon** and human **beta-interferon** have been used in this manner. (See, G. Gross, et al., "**Alpha-Interferon** in Condylomata Acuminata and Juvenile **Diabetes Mellitus**," Dtsch-Med.-Wochenschr, 1986, Sep. 5, III(36), pp. 1351-5; A. Schonfeld, et al., "Intramuscular Human **Interferon Beta** Injections in Treatment of Condylomata Acuminata," Lancet, 1984, May 12, I(8385), pp. 1038-42). Treatment of condylomata acuminata with interferon typically. . .

US PAT NO: 5,091,365 :IMAGE AVAILABLE:

L3: 6 of 6

DETDESC:

DETD(16)

insipidus

9

Corticotropin (ACTH)

inflammatory disorders

39

Tetracosactide

inflammatory disorders

24

Alsactide

"

17

Insulin

diabetes mellitus

51

.delta.-Sleep-ind.

sleep disturbances

9

peptide

Secretin

gastric hemorrhages

27

Cholecystokinin

biliary tract disorders,

8-32. . . III

cardiac and renal

24

insufficiency

ANF-(99-126)

cardiac and renal

28

insufficiency

Thymopentin

rheumatoid arthritis

5

Interferon-.alpha.

colds

125

Thyroliberin (TRH)

pituitary diagnostic aid

3

Gonadoliberin (LHRH)

cryptorchidism, sterility

10

Buserelin

prostate. . .

L1 ANSWER 33 OF 53 WPIDS COPYRIGHT 1998 DERWENT INFORMATION LTD
AN 94-302673 [37] WPIDS
DNC C94-159283

TI Use of alpha- or beta-**interferon** or analogues - for
preventing or treating an autoimmune disorder, e.g. **diabetes**
, arthritis, or transplant rejection.

DC B04 D16

IN SOBEL, D O

PA (GEOU) UNIV GEORGETOWN

CYC 18

PI WO 9420122 A1 940915 (9437)* 36 pp

RW: AT BE CH DE DK ES FR GB GR IE IT LU MC NL PT SE

W: AU CA

AU 9463549 A 940926 (9503)

ADT WO 9420122 A1 WO 94-US2154 940307; AU 9463549 A AU 94-63549 940307

FDT AU 9463549 A Based on WO 9420122

PRAI US 93-26758 930305

AB WO 9420122 A UPAB: 941223

A method of preventing or treating an autoimmune disease in a mammal
comprises administering at least one subtype of alpha- or beta-
interferon or a hybrid or analogue of either or a mixt. Also
claimed are:

(1) a method treating an asymptomatic preclinical autoimmune
state in a mammal, which comprises administering a single subtype of
alpha- or beta- **interferon** or a hybrid or analogue of
either or a mixt.; (1) a method inhibiting rejection of transplanted
islet cells or a pancreas in a mammal having transplanted islet
cells or pancreas, comprising administering a single subtype of
alpha- or beta-**interferon** or a hybrid or analogue or a
mixt.

USE - The method can be used for treating or preventing
autoimmune disorders such as type I **diabetes** mellitus,
rheumatoid arthritis, systemic lupus erythematosus, scleroderma,
sjogrens syndrome, mixed connective tissue disease, ankylosis
spondylitis, Reiter's syndrome, psoriatic arthritis,
hypersensitivity vasculitis, ulcerative colitis, cirrhosis,
autoimmune uveitis, myasthenia gravis, Buerger's disease, Kawasaki's
disease, systemic necrotising vasculitis, regional enteritis and
hypoparathyroidism.

The **interferon** can be administered at a dose of e.g.
1x105 units to 75x106 units, e.g. orally.

Dw

L6 ANSWER 600 OF 697 CAPLUS COPYRIGHT 1998 ACS DUPLICATE 249
AN 1989:93290 CAPLUS
DN 110:93290
TI Effect of interferons and poly(I):poly(C) on the pathogenesis of the
diabetogenic variant of encephalomyocarditis virus in different
mouse strains
AU Giron, David J.; Agostini, Heidi J.; Thomas, Donald C.
CS Coll. Sci. Math., Wright State Univ., Dayton, OH, USA
SO J. Interferon Res. (1988), 8(6), 745-53
CODEN: JIREDJ; ISSN: 0197-8357
DT Journal
LA English
AB **Interferon** (IFN) can either prevent or exacerbate the
pathogenic effects of the diabetogenic variant of
encephalomyocarditis (EMC-D) virus. The effect seen is dependent
upon the mouse strain and the time of IFN administration. Studies
were initiated to investigate the role of the IFN system in the
pathogenesis of this virus infection. Here IFNs or poly(I):poly(C)
were administered to several mouse strains at 24 h before or 4 days
after infection with EMC-D virus. The results of such treatment
ranged from complete protection of the animals from the diabetogenic
effects of the virus to exacerbation of the infection as reflected
by the virus content in selected organs. The effect was dependent
upon the mouse strain, the type of IFN, and the time of its
administration in relation to virus infection.

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L1 ANSWER 33 OF 53 WPIDS COPYRIGHT 1998 DERWENT INFORMATION LTD
 AN 94-302673 [37] WPIDS
 DNC C94-159283
 TI Use of alpha- or beta-**interferon** or analogues - for preventing or treating an autoimmune disorder, e.g. **diabetes**, arthritis, or transplant rejection.
 DC B04 D16
 IN SOBEL, D O
 PA (GEOU) UNIV GEORGETOWN
 CYC 18
 PI WO 9420122 A1 940915 (9437)* 36 pp
 RW: AT BE CH DE DK ES FR GB GR IE IT LU MC NL PT SE
 W: AU CA
 AU 9463549 A 940926 (9503)
 ADT WO 9420122 A1 WO 94-US2154 940307; AU 9463549 A AU 94-63549 940307
 FDT AU 9463549 A Based on WO 9420122
 PRAI US 93-26758 930305
 AB WO 9420122 A UPAB: 941223
 A method of preventing or treating an autoimmune disease in a mammal comprises administering at least one subtype of alpha- or beta-**interferon** or a hybrid or analogue of either or a mixt. Also claimed are:
 (1) a method treating an asymptomatic preclinical autoimmune state in a mammal, which comprises administering a single subtype of alpha- or beta- **interferon** or a hybrid or analogue of either or a mixt.; (1) a method inhibiting rejection of transplanted islet cells or a pancreas in a mammal having transplanted islet cells or pancreas, comprising administering a single subtype of alpha- or beta-**interferon** or a hybrid or analogue or a mixt.
 USE - The method can be used for treating or preventing autoimmune disorders such as type I **diabetes** mellitus, rheumatoid arthritis, systemic lupus erythematosus, scleroderma, sjogrens syndrome, mixed connective tissue disease, ankylosis spondylitis, Reiter's syndrome, psoriatic arthritis, hypersensitivity vasculitis, ulcerative colitis, cirrhosis, autoimmune uveitis, myasthenia gravis, Buerger's disease, Kawasaki's disease, systemic necrotising vasculitis, regional enteritis and hypoparathyroidism.
 The **interferon** can be administered at a dose of e.g. 1x10⁵ units to 75x10⁶ units, e.g. orally.
 Dw

AN 86300315 MEDLINE
DN 86300315
TI [Alpha **interferon** in **condylomata** acuminata and
juvenile diabetes mellitus].
Interferon-alpha bei **Condylomata** acuminata und
juvenilem Diabetes mellitus.
AU Gross G; Roussaki A; Ikenberg H; Drees N
SO DEUTSCHE MEDIZINISCHE WOCHENSCHRIFT, (1986 Sep 5) 111 (36) 1351-5.
Journal code: ECL. ISSN: 0012-0472.
CY GERMANY, WEST: Germany, Federal Republic of
DT Journal; Article; (JOURNAL ARTICLE)
LA German
FS Priority Journals; Cancer Journals
EM 198612
AB Persistent condylomata acuminata in a 21-year-old patient with
diabetes mellitus were treated with highly purified interferon-alpha
(IFN-alpha) obtained by recombinant DNA technology. Daily dose was
1.5 X 10(6) IU, given subcutaneously. Already during treatment the
condylomata regressed. Two weeks after the end of therapy, i.e.
after a total dose of 10.5 X 10(6) IU IFN-alpha, all condylomata had
completely receded. Blood glucose levels remained constant with
concomitant insulin therapy. Toxic side-effects or antibodies to
IFN-alpha were not observed.